

June 2, 2025

BSE Limited

Code: 532321

1st Floor,
P J Towers,
Dalal Street,
Mumbai-400001

National Stock Exchange of India Limited

Code: Zyduslife

Exchange Plaza, 5th Floor,
Plot No. C/1, G Block,
Bandra-Kurla Complex, Bandra (East),
Mumbai-400051

Re.: Press Release

Dear Sir / Madam,

Please find enclosed a copy of press release dated June 2, 2025, titled "Zydus receives tentative approval from USFDA for Rifaximin Tablets, 550 mg."

The contents of the press release give full details.

Please bring the aforesaid news to the notice of the members of the exchange and the investors' at large.

Thanking you,

Yours faithfully,

For, **ZYDUS LIFESCIENCES LIMITED**

DHAVAL N. SONI

COMPANY SECRETARY AND COMPLIANCE OFFICER

MEMBERSHIP NO. FCS7063

Encl.: As above



Zydus receives tentative approval from USFDA for Rifaximin Tablets, 550 mg

Ahmedabad, India, June 2, 2025

Zydus Lifesciences Limited (including its subsidiaries/ affiliates, hereafter referred to as “Zydus”) has received tentative approval from the United States Food and Drug Administration (USFDA) for Rifaximin Tablets, 550 mg (USRLD: Xifaxan® Tablets, 550 mg).

Rifaximin tablets are indicated for the treatment of irritable bowel syndrome with diarrhoea (IBS-D) in adults. Rifaximin tablets will be produced at the Group’s manufacturing site at SEZ II, Ahmedabad.

Rifaximin tablets had annual sales of USD \$2672.9 mn in the United States (IQVIA MAT March 2025).

The group now has 427 approvals and has so far filed 492* ANDAs since the commencement of the filing process in FY 2003-04.

*(*As on 31st March, 2025)*



**PRESS
RELEASE**

For further information please contact :
The Corporate Communications Department

Zydus Lifesciences Limited

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